TÜVRheinland®

EC Certificate

Directive 93/42/EEC Annex V
Production Quality Assurance
Medical Devices

Registration No.: DD 60148109 0001

Report No.: 50180180 006

Manufacturer:
Lyncmed Medical Technology
(Beijing) Co., Ltd.
Room 1601, Building No.2
Zhubang 2000 Business Building, Balizhuangxili 99
Chaoyang District
100022 Beijing
P.R. China

Products: Sterile Latex Surgical Gloves

Replaces Approval, Registration No.: DD 60133385 0001

Expiry Date: 2023-12-03

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Effective Date: 2020-06-22

Date: 2020-06-22

Notified Body
TÜV Rheinland LGA Products GmbH
Zertifizierstelle
Jing Zhang

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

Digitally signed by Perminov Andrei
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